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NEWS NEWS	1 2	NOV	21	Web Page for STN Seminar Schedule - N. America CAS patent coverage to include exemplified prophetic substances identified in English-, French-, German-,
MEGG	2	MOTE	0.0	and Japanese-language basic patents from 2004-present
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				coverage of complete UK patent families
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NEWS	10	JAN	07	WPIDS, WPINDEX, and WPIX enhanced Japanese Patent Classification Data
NEWS	11	FEB	02	Simultaneous left and right truncation (SLART) added for CERAB, COMPUAB, ELCOM, and SOLIDSTATE
NEWS	12	FEB	0.2	GENBANK enhanced with SET PLURALS and SET SPELLING
NEWS		FEB		Patent sequence location (PSL) data added to USGENE
NEWS		FEB		COMPENDEX reloaded and enhanced
NEWS		FEB		WTEXTILES reloaded and enhanced
NEWS		FEB		New patent-examiner citations in 300,000 CA/CAplus
NEWS	10	CLD	13	patent records provide insights into related prior art
NEWS	17	FEB	19	Increase the precision of your patent queries use terms from the IPC Thesaurus, Version 2009.01
NEWS	18	FEB	23	Several formats for image display and print options discontinued in USPATFULL and USPAT2
NEWS	19	FEB	23	MEDLINE now offers more precise author group fields
NEWS	20	FEB	23	and 2009 MeSH terms TOXCENTER updates mirror those of MEDLINE - more
				precise author group fields and 2009 MeSH terms
NEWS	21	FEB	23	Three million new patent records blast AEROSPACE into STN patent clusters
NEWS	22	FEB	25	USGENE enhanced with patent family and legal status display data from INPADOCDB
NEWS	23	MAR	06	INPADOCDB and INPAFAMDB enhanced with new display formats
NEWS	24	MAR	11	EPFULL backfile enhanced with additional full-text applications and grants
NEWS	25	MAR	11	ESBIOBASE reloaded and enhanced
NEWS		MAR		CAS databases on STN enhanced with new super role
NEMS	20	HHI	20	for nanomaterial substances
NEWS	27	MAR	23	CA/CAplus enhanced with more than 250,000 patent equivalents from China
				-1

NEWS EXPRESS JUNE 27 08 CURRENT WINDOWS VERSION IS V8.3, AND CURRENT DISCOVER FILE IS DATED 23 JUNE 2008.

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=> index bioscience medicine FILE 'DRUGMONOG' ACCESS NOT AUTHORIZED COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.22 0.22

FULL ESTIMATED COST

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, ANTE, AQUALINE, AQUASCI, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CAPLUS, CEABA-VTB, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DISSABS, DRUGB, DRUGMONOG2, DRUGU, EMBAL, EMBASE, ...' ENTERED AT 17:20:47 ON 26 MAR 2009

71 FILES IN THE FILE LIST IN STNINDEX

Enter SET DETAIL ON to see search term postings or to view search error messages that display as 0\* with SET DETAIL OFF.

- => s (asparaginas? or (asparagin?(3a)amidohydrolas?))
  - 476 FILE ADISCTI
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  - 135 FILE ADISNEWS
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  - 4307 FILE CAPLUS
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1471

USPAT2

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F63	1	NUTRACEUT
F64	1	PS
F65	1	WPIFV
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PROCESSING COMPLETED FOR L4
L5 31 DUP REM L4 (11 DUPLICATES REMOVED)

=> d ti 15 1-31

L5 ANSWER 1 OF 31 USPATFULL on STN TI Process for Reducing Acrylamide

L5 ANSWER 2 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 1

TI Aspergillus niger asparaginase variants and their commercial uses

L5 ANSWER 3 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 2

- TI Aspergillus niger asparaginase variants and their commercial uses
- L5 ANSWER 4 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 3
- TI Design of thermostable asparaginases and their use for reduction of acrylamide in foods
- L5 ANSWER 5 OF 31 USPATFULL on STN
- TI Process Flavours with Low Acrylamide
- L5 ANSWER 6 OF 31 USPATFULL on STN
- TI PROCESS FOR TREATING VEGETABLE MATERIAL WITH AN ENZYME
- L5 ANSWER 7 OF 31 USPATFULL on STN
- TI Methods for reducing asparagine in a food material using cooling
- L5 ANSWER 8 OF 31 USPATFULL on STN
- TI Methods for reducing asparagine in a dough food component using water activity
- L5 ANSWER 9 OF 31 USPATFULL on STN
- TI Method of Preparing a Heat-Treated Product
- L5 ANSWER 10 OF 31 USPATFULL on STN
- TI Asparaginases and Method of Preparing a Heat-Treated Product
- L5 ANSWER 11 OF 31 USPATFULL on STN
- TI Amidases from Aspergillus Niger and Their Use in a Food Production Process
- L5 ANSWER 12 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TI Production of L-asparaginase by isolated Aspergillus species using SSF
- L5 ANSWER 13 OF 31 USPATFULL on STN
- TI Novel food production process
- L5 ANSWER 14 OF 31 USPATFULL on STN
- TI Novel food production process
- L5 ANSWER 15 OF 31 USPATFULL on STN
- TI Functionalization of yarn and textile products
- L5 ANSWER 16 OF 31 SCISEARCH COPYRIGHT (c) 2009 The Thomson Corporation on STN
- TI Development and application of aspergillus niger asparaginase to prevent the formation of acrylamide in food products
- L5 ANSWER 17 OF 31 USPATFULL on STN DUPLICATE 4
- TI Method of preparing a heat-treated product
- L5 ANSWER 18 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- ${\tt TI}$  Detection of the antitumor glutaminase-asparaginase in the filamentous fungi
- L5 ANSWER 19 OF 31 EMBASE COPYRIGHT (c) 2009 Elsevier B.V. All rights reserved on STN DUPLICATE 5
- TI Production of L-asparaginase, an anticancer agent, from Aspergillus niger using agricultural waste in solid state fermentation.
- L5 ANSWER 20 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN

- TI Enzymic processing of food to limit acrylamide formation
- L5 ANSWER 21 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 6
- TI Food production process involving asparaginase yielded from recombinant Aspergillus niger
- L5 ANSWER 22 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 7
- TI Purification and properties of L-asparaginase produced by Aspergillus niger, S-48 TAT, the causal fungus of biodeterioration inside Tut Ankhamen Tomb (TAT)
- L5 ANSWER 23 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TI Factors affecting the production and activity of fungal asparaginases using whey
- L5 ANSWER 24 OF 31 BIOSIS COPYRIGHT (c) 2009 The Thomson Corporation on STN
- TI ENZYMES IMMOBILIZED ON ALUMINA AND STAINLESS STEEL SUPPORTS.
- L5 ANSWER 25 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TI Are the urease and asparaginase of Aspergillus niger endocellular enzymes?
- L5 ANSWER 26 OF 31 BIOSIS COPYRIGHT (c) 2009 The Thomson Corporation on STN
- TI L'urease et l'asparaginase de l'Aspergillus niger sont-elles des endo-diastases?.
- L5 ANSWER 27 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TI The evolution of urease in cultures of Aspergillus niger
- L5 ANSWER 28 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TI The evolution of asparaginase in cultures of Aspergillus niger
- L5 ANSWER 29 OF 31 BIOSIS COPYRIGHT (c) 2009 The Thomson Corporation on STN
- TI L'evolution de l'asparaginase dans les cultures de l' Aspergillus niger.
- L5 ANSWER 30 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TI Fermentative hydrolysis of asparagine by the Mycelium of Aspergillus niger
- L5 ANSWER 31 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TI Conditions of action of asparaginase of Aspergillus niger
- => d ibib abs 15 2 4 12 16 19 21 22 25 28 31
- L5 ANSWER 2 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 1

ACCESSION NUMBER: 2008:333054 TOXCENTER COPYRIGHT: Copyright 2009 ACS

DOCUMENT NUMBER: CA14923507705E

TITLE: Aspergillus niger asparaginase variants and their commercial uses

AUTHOR(S): Laan, Van Der Jan Metske; Stor, Mark Cristiaan; Lange, De

Ilse; Mohrmann, Lisette

CORPORATE SOURCE: ASSIGNEE: DSM IP Assets B. V. PATENT INFORMATION: WO 2008128975 A1 30 Oct 2008 SOURCE: (2008) PCT Int. Appl., 70pp.

CODEN: PIXXD2.

COUNTRY: NETHERLANDS

DOCUMENT TYPE: Patent FILE SEGMENT: CAPLUS

OTHER SOURCE: CAPLUS 2008:1299788

LANGUAGE: English

ENTRY DATE: Entered STN: 11 Nov 2008

Last Updated on STN: 2 Dec 2008

AB The present provides two polypeptide variants of Aspergillus niger asparaginase and to polynucleotide sequences that encode such novel asparaginase variants. The variants display a higher specific activity at the same pH, a higher pH optimum and broader pH-activity profile, and improved thermostability, in comparison to the wild-type enzyme. Furthermore, the invention relates to the use of these novel asparaginase variants in industrial processes, including the reduction of acrylamide formed in thermally processed food products via the maillard reaction and use a medicament in the treatment of tumors.

L5 ANSWER 4 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 3

ACCESSION NUMBER: 2008:308413 TOXCENTER COPYRIGHT: Copyright 2009 ACS CA14918396649T

TITLE: Design of thermostable asparaginases and their use for

reduction of acrylamide in foods

AUTHOR(S): Matsui, Tomoko; Friis, Esben Peter; Yamaqishi, Akihiko

CORPORATE SOURCE: ASSIGNEE: Novozymes A/S PATENT INFORMATION: WO 2008110513 A1 18 Sep 2008 SOURCE: (2008) PCT Int. Appl., 63pp.

CODEN: PIXXD2.

COUNTRY: DENMARK
DOCUMENT TYPE: Patent
FILE SEGMENT: CAPLUS

OTHER SOURCE: CAPLUS 2008:1119491

LANGUAGE: English

ENTRY DATE: Entered STN: 14 Oct 2008

Last Updated on STN: 4 Nov 2008

AΒ The invention relates to new asparaginases having improved properties, preferably improved thermotolerance, such as improved activity at high temps. and/or improved thermostability. The three-dimensional of an asparaginase from Aspergillus oryzae was modeled based on the published structure of a homologous enzyme from Erwinia chrysanthemi. Based on the modeled structure, amino acid residues are identified of relevance for improving the properties of the asparaginase, especially the thermotolerance. Further, an inferred ancestral asparaginase sequence was predicted, and from this sequence further amino acid residues of relevance for improving the properties of the asparaginase are identified. Based on such structural and functional considerations, asparaginase variants are constructed having modified amino acid residues at the identified positions and having altered physiochem. properties, especially improved relative activity at high temps. and/or improved thermostability. invention also relates to DNA sequences encoding such improved asparaginases, their production in a recombinant host cell, as well as methods of using the asparaginases, in particular for reduction of acrylamide in foods.

L5 ANSWER 12 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:1170798 CAPLUS

TITLE: Production of L-asparaginase by isolated Aspergillus

species using SSF

AUTHOR(S): Uppuluri, K. B.; Kalidindi, S. V.; Sekhar, P. V. G.

V.; Harish, Ch.; Reddy, D. S. Rami

CORPORATE SOURCE: Department of Biotechnology, Bapatla Engineering

College, Bapatla, 52101, India

SOURCE: Biosciences, Biotechnology Research Asia (2008), 5(1),

229-236

CODEN: BBRAB4; ISSN: 0973-1245

PUBLISHER: Oriental Scientific Publishing Co.

DOCUMENT TYPE: Journal LANGUAGE: English

AB Acute lymphocytic leukemia is a common leukemia characterized by frequent infections and anemia. Thousands of new cases are diagnosed each year worldwide. L-asparaginase (E.C.3.5.1.1), also known as L-asparagine amino hydrolase, is a potential anti tumor enzyme that catalyzes the hydrolysis of L-asparagine into L-aspartic acid and ammonia. L-asparaginase production was investigated in the filamentous fungi on sesame cake using solid state

fermentation(SSF). One-factor-at-a-time approach design was applied to

optimize

a solid-state fermentation using the sesame cake as a main substrate, for the production of L-asparaginase by isolated Aspergillus Species. Effect of Various environmental factors like Particle size of the solid medium, Moisture content, Incubation pH, Time and temperature and number of different nutritional supplements were verified on the activity and specific activity of extracellular enzyme, L-asparaginase. Among those pH, Particle size, moisture content, glucose, Ammonium sulfate and L-asparagine were the most significant factors improving enzyme production process. The second optimization step was carried out to identify the different sources of the three factors influencing the production of enzyme namely Glucose, ammonium sulfate and L-asparagine, that bringing about maximum L-asparaginase activity. Maximal enzyme activity (191.2 IU) has been detected under the following conditions, pH 6.5, temperature 32°C, incubation period 108 h, particle size of 0.67 cm, moisture content of 1:1 (Media: buffer) when medium was supplemented with 3%weight/weight Fructose, 3%weight/weight Ammonium sulfate, 0.1%weight/weight L-Asparagine, 0.01% weight/weight

Magnesium Sulfate, 01% weight/weight sodium chloride and inoculum size of 1.5ml (1.6 + 103 Spores/mL) which is nearly three folds the activity in

basal medium.

REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 16 OF 31 SCISEARCH COPYRIGHT (c) 2009 The Thomson Corporation on

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ACCESSION NUMBER: 2007:953935 SCISEARCH

THE GENUINE ARTICLE: 192FE

TITLE: Development and application of aspergillus

niger asparaginase to prevent the

formation of acrylamide in food products

AUTHOR: Koster, F.

CORPORATE SOURCE: DSM Food Specialties, Delft, Netherlands

COUNTRY OF AUTHOR: Netherlands

SOURCE: ANNALS OF NUTRITION AND METABOLISM, (2007) Vol. 51, Supp.

[1], pp. 393-393. ISSN: 0250-6807.

PUBLISHER: KARGER, ALLSCHWILERSTRASSE 10, CH-4009 BASEL, SWITZERLAND.

DOCUMENT TYPE: Conference; Journal

LANGUAGE: English

REFERENCE COUNT: 0

ENTRY DATE: Entered STN: 4 Oct 2007

Last Updated on STN: 4 Oct 2007

L5 ANSWER 19 OF 31 EMBASE COPYRIGHT (c) 2009 Elsevier B.V. All rights reserved on STN DUPLICATE 5

ACCESSION NUMBER: 2006520364 EMBASE

TITLE: Production of L-asparaginase, an anticancer

agent, from Aspergillus niger using

agricultural waste in solid state fermentation.

AUTHOR: Mishra, Abha (correspondence)

CORPORATE SOURCE: School of Biochemical Engineering, Institute of Technology,

Banaras Hindu University, Varanasi-221005, India.

abha91@yahoo.co.in

SOURCE: Applied Biochemistry and Biotechnology, (Oct 2006) Vol.

135, No. 1, pp. 33-42.

Refs: 20

ISSN: 0273-2289

PUBLISHER IDENT: ABAB135133
COUNTRY: United States
DOCUMENT TYPE: Journal; Article

FILE SEGMENT: 037 Drug Literature Index

004 Microbiology: Bacteriology, Mycology, Parasitology

and Virology

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 10 Nov 2006

Last Updated on STN: 10 Nov 2006

AΒ This article reports the production of high levels of Lasparaginase from a new isolate of Aspergillus niger in solid state fermentation (SSF) using agrowastes from three leguminous crops (bran of Cajanus cajan, Phaseolus mungo, and Glycine max). When used as the sole source for growth in SSF, bran of G. max showed maximum enzyme production followed by that of P. mungo and C. cajan. A 96-h fermentation time under aerobic condition with moisture content of 70%, 30 min of cooking time and 1205-1405  $\mu$  range of particle size in SSF appeared optimal for enzyme production. Enzyme yield was maximum (40.9  $\pm$  3.35 U/g of dry substrate) at pH 6.5 and temperature 30  $\pm$  2°C. The optimum temperature and pH for enzyme activity were 40°C and 6.5, respectively. The study suggests that choosing an appropriate substrate when coupled with process level optimization improves enzyme production markedly. Developing an asparaginase production process based on bran of G. max as a substrate in SSF is economically attractive as it is a cheap and readily available raw material in agriculture-based countries. Copyright .COPYRGT. 2006 by

L5 ANSWER 21 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 6

ACCESSION NUMBER: 2004:102087 TOXCENTER COPYRIGHT: Copyright 2009 ACS DOCUMENT NUMBER: CA14021338277Y

TITLE: Food production process involving asparaginase

yielded from recombinant Aspergillus

niger

AUTHOR(S): Plomp, Pieter Jan Arnoldus Maria; De Boer, Lex; Van

Rooijen, Rutger Jan; Meima, Roelf Bernhard

CORPORATE SOURCE: ASSIGNEE: DSM Ip Assets B.V. PATENT INFORMATION: WO 2004030468 A2 15 Apr 2004 SOURCE: (2004) PCT Int. Appl., 46 pp.

CODEN: PIXXD2.

COUNTRY: NETHERLANDS

DOCUMENT TYPE: Patent FILE SEGMENT: CAPLUS

Humana Press Inc.

OTHER SOURCE: CAPLUS 2004:308353

LANGUAGE: English

ENTRY DATE: Entered STN: 4 May 2004

Last Updated on STN: 22 Jan 2008

AB A process for the production of a food product involving at least one heating step, comprises adding one or more enzymes to an intermediate form of the food product in the production process. The enzyme is added prior to the heating step in an amount that is effective in reducing the level of amino

acids that are present in the intermediate form of the food product which amino acids are involved in the formation of acrylamide during the heating step. The invention also relates to food products obtained from the process. Thus, the asparaginase encoded by a nucleotide sequence is obtained by constructing expression plamids containing the DNA sequence, transforming an Aspergillus niger strain with this plamid, and growing the strain.

L5 ANSWER 22 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 7

ACCESSION NUMBER: 1999:196080 TOXCENTER COPYRIGHT: Copyright 2009 ACS CA13126348273X

TITLE: Purification and properties of L-asparaginase

produced by Aspergillus niger, S-48

TAT, the causal fungus of biodeterioration inside Tut

Ankhamen Tomb (TAT)

AUTHOR(S): Louboudy, S. S.

CORPORATE SOURCE: Bot. & Microbiol, Dept., Fac. of Sci., Al-Azhar Univ.,

Cairo, Egypt.

SOURCE: Egyptian Journal of Biotechnology, (1998) Vol. 4, pp.

110-123.

CODEN: EJBIF7. ISSN: 1110-6093.

COUNTRY: EGYPT
DOCUMENT TYPE: Journal
FILE SEGMENT: CAPLUS

OTHER SOURCE: CAPLUS 1999:649978

LANGUAGE: English

ENTRY DATE: Entered STN: 16 Nov 2001

Last Updated on STN: 9 May 2002

AB The purification and properties of L-asparaginase (I) produced by A. niger S-48 TAT, the causal factor of biodeterioration inside the Pharaoh Tutankhamen tomb (TAT), is reported. The purification procedure involved cell-free filtrate preparation (specific activity of 8.92 U/mg protein/mL), fractional precipitation with (NH4)2SO4, (specific activity of 21.05

U/mg protein/mL corresponding to a 2.35-fold purification), dialysis against distilled water followed by dialysis against sucrose crystals, (specific activity of 36.92 U/mg protein/mL, corresponding to a 5.7-fold purification) and finally applying a column of Sephadex G-100 (specific activity of 61.0 U/mg protein/mL corresponding to a 6.83-fold purification). The regulatory role of different buffers applied at different pH values revealed that purified I exhibited a maximum specific activity of 62.8 U/mg protein/mL in the presence of citrate-phosphate buffer pH 6.6, followed by citrate buffer pH 6.0 (specific activity of 55.46 U/mg protein/mL) and then Tris-HCl buffer pH 7.4 which revealed an obvious decrease in the specific activity (34.16 U/mg protein/mL). By testing purified I in the presence of different substrates, it was found that the highest activity was obtained by using the most preferable one, i.e., L-asparagine, followed by L-aspartic acid, L-glutamine, and L-glutamic acid, whereas L-arginine, L-ornithine, L-threonine and L-citrulline showed negligible or inhibitory effects toward the purified enzyme activity. Moreover, the application of different heavy metal cations (in the form of chloride salts in addition to KCN) as activators and/or inhibitors indicated that CaCl2, NH4Cl, BaCl2, and MnCl2 promoted I activity, whereas AlCl3, KCN, NiCl2, ZnCl2, FeCl2, and MgCl2 had deleterious effects on enzyme activity. Purified I was tested at different incubation temps., and showed obvious activity within the temperature range of  $22.5-45^{\circ}$  with a maximum at  $30^{\circ}$ .

L5 ANSWER 25 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1930:23193 CAPLUS

DOCUMENT NUMBER: 24:23193 ORIGINAL REFERENCE NO.: 24:2478g-h TITLE: Are the urease and asparaginase of

Aspergillus niger endocellular

enzymes?

AUTHOR(S): Bach, D.

SOURCE: Bulletin de la Societe de Chimie Biologique (1929),

11, 1016-24

CODEN: BSCIA3; ISSN: 0037-9042

DOCUMENT TYPE: Journal LANGUAGE: Unavailable

AB The dried and finely ground mycelium of Aspergillus

niger lost about 2/3 of its urease and asparaginase

activity when suspended in buffer solution and filtered through paper. If itrate from a Chamberland filter was practically inactive. Enzyme activity was also greatly reduced by long-continued maceration. It is concluded from these and other expts. described previously that both

enzymes are endocellular.

L5 ANSWER 28 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1930:23191 CAPLUS

DOCUMENT NUMBER: 24:23191
ORIGINAL REFERENCE NO.: 24:2478e-f

TITLE: The evolution of asparaginase in cultures of

Aspergillus niger

AUTHOR(S): Bach, D.

SOURCE: Bulletin de la Societe de Chimie Biologique (1929),

11, 995-1006

CODEN: BSCIA3; ISSN: 0037-9042

DOCUMENT TYPE: Journal LANGUAGE: Unavailable

AB cf. C. A. 24, 1133. Asparaginase is an endocellular enzyme which is normally present independently of asparagine in the media. The amount present declines to a min. in 6 days, rises to a maximum in 10 days, and steadily declines to 20 days. The asparaginase activity is parallel with the general proteolytic activity for supplying NH3 to the cultures.

L5 ANSWER 31 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 1929:7263 CAPLUS

DOCUMENT NUMBER: 23:7263
ORIGINAL REFERENCE NO.: 23:861h-i

TITLE: Conditions of action of asparaginase of

Aspergillus niger

AUTHOR(S): Bach, D.

SOURCE: Compt. rend. (1928), 187, 955-6

DOCUMENT TYPE: Journal LANGUAGE: Unavailable

The enzyme was active only in neutral or slightly alkaline media, the optimum PH being 84 to 88. At the same time Aspergillus niger was able to utilize asparagine completely in media more acid than pH 6.4. The optimum temperature varied with the pH of the medium being 42° for pH 8.6 and 31° for pH 7.6. The temperature zone of action was wide, extending from 7° to 70°. When the concentration of the substrate (asparagine) increased beyond 1%, the NH3 produced tended toward a limit which was independent of the concentration of the substrate. Complete hydrolysis of the asparagine was not attained, but reached about 80% under optimum conditions. As the hydrolysis proceeded there was a diminution in its velocity after about 36 hrs., due principally to a destruction of the enzyme. The presence of asparagine tended to protect the asparaginase from autodestruction.

- L5ANSWER 22 OF 31 TOXCENTER COPYRIGHT 2009 ACS on STN DUPLICATE 7
- ТΤ Purification and properties of L-asparaginase produced by Aspergillus niger, S-48 TAT, the causal fungus of biodeterioration inside Tut Ankhamen Tomb (TAT)
- The purification and properties of L-asparaginase (I) produced by A. AB niger S-48 TAT, the causal factor of biodeterioration inside the Pharaoh Tutankhamen tomb (TAT), is reported. The purification procedure involved cell-free.
- L5ANSWER 31 OF 31 CAPLUS COPYRIGHT 2009 ACS on STN
- TIConditions of action of asparaginase of Aspergillus niger
- IT Aspergillus niger

(asparaginase of, conditions of action of)

- IT 70-47-3, Asparagine
  - (hydrolysis by asparaginase of Aspergillus niger)
- ΤТ 9015-68-3, Asparaginase

(in Aspergillus niger, conditions of action of)

=> FIL STNGUIDE

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 92.34 94.60 DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE -3.28-3.28

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FILE USPATFULL
FILE COVERS 1971 TO PATENT PUBLICATION DATE: 26 Mar 2009 (20090326/PD)
FILE LAST UPDATED: 26 Mar 2009 (20090326/ED)
HIGHEST GRANTED PATENT NUMBER: US7509687
HIGHEST APPLICATION PUBLICATION NUMBER: US20090083889
CA INDEXING IS CURRENT THROUGH 26 Mar 2009 (20090326/UPCA)
ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 26 Mar 2009 (20090326/PD)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2008
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Dec 2008

USPATFULL now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

## FILE EMBASE

FILE COVERS 1974 TO 26 Mar 2009 (20090326/ED)

EMBASE was reloaded on March 30, 2008.

EMBASE is now updated daily. SDI frequency remains weekly (default) and biweekly.

This file contains CAS Registry Numbers for easy and accurate substance identification.

Beginning January 2008, Elsevier will no longer provide EMTREE codes as part of the EMTREE thesaurus in EMBASE. Please update your current-awareness alerts (SDIs) if they contain EMTREE codes.

For further assistance, please contact your local helpdesk.

FILE TOXCENTER

FILE COVERS 1907 TO 24 Mar 2009 (20090324/ED)

The MEDLINE file segment has been reload and updated with the National Library of Medicine's revised 2009 MeSH terms. See HELP RLOAD for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

The BIOSIS segment of TOXCENTER has been augmented with 13,000 records from 1946 through 1968.

FILE CAPLUS

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